



K014147

		<b>SPECIAL 510(k) 80 CHANNEL EEG</b>
<b>SPECIAL 510(k) DEVICE MODIFICATION</b>		<b>DECEMBER 14, 2001</b> PAGE 49 of 51

## Section F – 510(k) SUMMARY

This summary of 510(k) safety and effectiveness information is being submitted in accordance with the requirements of the SMDA 1990 and 21 CFR § 807.92.

**Name:** Cameron Mahon **JAN 14 2002**  
Vice President, Customer Satisfaction

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Canada, L6H 5S1

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**Common Names:** 80 Channel EEG

**Classification Name:** Electroencephalograph

**Predicate Devices:** 24 Channel Ambulatory EEG [FDA 510(k) K982479]

**Description:** The 80 Channel EEG is a digital electroencephalograph

**Substantial Equivalence:** The 80 Channel EEG is substantially equivalent to the 24 Channel Ambulatory EEG [FDA 510(k) K982479]

**Indications for Use:** The 80 Channel EEG is intended to be used as an electroencephalograph: to acquire, display, store, and archive electroencephalographic signals.



Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20850

**JAN 14 2002**

XLTEK  
Sonja Markez  
Regulatory Affairs  
2568 Bristol Circle  
Oakville, Ontario  
Canada L6H 5S1

Re: K014147  
Trade Name: 80 Channel EEG  
Regulation Number: 882.1400  
Regulation Name: Electroencephalograph  
Regulatory Class: II  
Product Code: GWQ  
Dated: December 14, 2001  
Received: December 18, 2001

Dear Ms. Markez:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

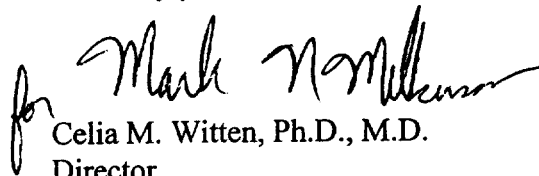
If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 21 CFR Part 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4659. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its Internet address <http://www.fda.gov/cdrh/dsma/dsmamain.html>

Sincerely yours,

for Mark N. Miller

Celia M. Witten, Ph.D., M.D.

Director



Division of General, Restorative  
and Neurological Devices

Office of Device Evaluation

Center for Devices and

Radiological Health

Enclosure

		<i>SPECIAL 510(k)</i> <i>80 CHANNEL EEG</i>
<i>SPECIAL 510(k) DEVICE MODIFICATION</i>		<i>DECEMBER 14, 2001</i> <i>PAGE 50 of 51</i>

## Section G – INDICATIONS FOR USE

510(k) Number (if known): K014141

Device Name: 80 Channel EEG

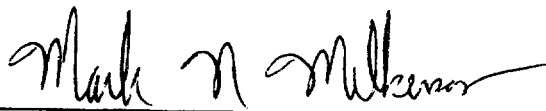
Indications for Use: The 80 Channel EEG is intended to be used as an electroencephalograph: to acquire, display, store, and archive electroencephalographic signals.

(PLEASE DO NOT WRITE BELOW THIS LINE – CONTINUE ON ANOTHER PAGE IF NEEDED)

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Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use X OR Over-The Counter Use \_\_\_\_\_  
(Per 21§ CFR 801.109)

for   
(Division Sign-Off)

Division of General, Restorative  
and Neurological Devices

(Optional Format 1-2-96)

510(k) Number K014147